

**Pharmacy Prior Authorization  
Non-Formulary, Step Therapy and Prior Authorization Guidelines**

Scroll down to see PA Criteria by drug class, or Ctrl+F to each document by drug name

PA Guideline	Requirements	Duration of Approval if Requirements Are Met
<p><b>Corlanor<sup>i</sup></b></p>	<p><b>May be authorized for members 18 years of age or older when the following criteria are met:</b></p> <ul style="list-style-type: none"> <li>• Diagnosis of stable symptomatic chronic heart failure (New York Heart Association (NYHA) Class II-III)</li> <li>• Left ventricular ejection fraction (LVEF) is less than or equal to 35%</li> <li>• Member is in sinus rhythm with a resting heart rate greater than or equal to 70 beats per minute</li> <li>• Continuation of therapy with maximally tolerated beta-blocker, or there is intolerance or contraindication to beta-blockers</li> <li>• Continuation of therapy with angiotensin-converting-enzyme inhibitor (ACEI)/Angiotensin Receptor Blockers (ARB), or Entresto, or there is intolerance, or contraindication to angiotensin-converting-enzyme inhibitor (ACEI)/Angiotensin Receptor Blockers (ARB), or Entresto</li> <li>• Provider attestation that no contraindications to treatment exist:                             <ul style="list-style-type: none"> <li>○ Acute decompensated heart failure</li> <li>○ Blood pressure less than 90/50 mmHg</li> <li>○ Pacemaker dependent (for example: heart rate maintained exclusively by pacemaker)</li> <li>○ Sick sinus syndrome, sinoatrial block of third-degree AV block (unless functioning</li> </ul> </li> </ul>	<p><b>Initial Approval:</b> 6 months</p> <p><b>Renewals:</b> 1 year</p> <p><b>Requires:</b></p> <ul style="list-style-type: none"> <li>• Member is responding to treatment</li> <li>• Heart rate is within recommended range for continuation of maintenance dose                             <ul style="list-style-type: none"> <li>• For example, 50-60 beats per minute, or dose adjusted</li> </ul> </li> </ul>

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	<ul style="list-style-type: none"> <li>○ demand pacemaker is present)</li> <li>○ Severe hepatic impairment (Child-Pugh class C)</li> </ul> <p><b>May be authorized for pediatric members 6 months of age or older when the following criteria are met:</b></p> <ul style="list-style-type: none"> <li>• Diagnosis of heart failure due to dilated cardiomyopathy</li> <li>• Member is in sinus rhythm with a resting heart rate of greater than or equal to 70 beats per minute</li> <li>• Provider attestation that no contraindications to treatment exist:                             <ul style="list-style-type: none"> <li>○ Acute decompensated heart failure</li> <li>○ Blood pressure less than 90/50 mmHg</li> <li>○ Pacemaker dependent (for example, heart rate maintained exclusively by pacemaker)</li> <li>○ Sick sinus syndrome, sinoatrial block or third-degree AV block (unless functioning demand pacemaker is present)</li> <li>○ Severe hepatic impairment (Child-Pugh class C)</li> </ul> </li> </ul>	<p>accordingly to achieve goal</p> <p><b>Quantity Level Limit:</b> Adults and Pediatrics: 60 tablets per 30 days</p> <p>Oral solution for pediatrics: 120 ampules per 30 days</p>
<b>Egrifta SV<sup>ii</sup></b>	<ul style="list-style-type: none"> <li>• Diagnosis of human immunodeficiency virus (HIV)-associated lipodystrophy</li> <li>• Prescribed by or in consultation with an infectious disease specialist</li> </ul>	<p><b>Initial Approval:</b> 6 months</p>

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	<ul style="list-style-type: none"> <li>• Documentation of waist circumference greater than or equal to 95 cm for males, or greater than or equal to 94 cm for females at start of therapy</li> <li>• Member is currently receiving and stable on anti-retroviral therapy for at least 8 weeks</li> <li>• Member does not have an active malignancy</li> <li>• Member does not have disruption of the hypothalamic-pituitary gland axis, or head trauma</li> <li>• Women of childbearing potential are not pregnant and are using appropriate contraception</li> <li>• Egrifta will not be approved for weight loss management</li> </ul>	<p><b>Renewal Approval:</b> 6 months</p> <p><b>Requires:</b></p> <ul style="list-style-type: none"> <li>• Documentation of a positive clinical response to treatment: A decrease in waist circumference</li> </ul>

**<sup>1</sup> Corlanor References**

1. Yancy CW et al. 2017 ACC/AHA/HFSA Focused Update of the 2013 ACCF/AHA Guideline for the Management of Heart Failure Circulation: 2017. <http://www.onlinejacc.org/content/accj/70/6/776.full.pdf? ga=2.179733604.1964533065.1574204551-936785029.1560984365>. Accessed November 19, 2019.
2. Corlanor (ivabradine) [package insert]. Thousand Oaks, CA; Amgen Inc.; Revised April, 2019. Retrieved from [https://www.pi.amgen.com/~media/amgen/repositorysites/pi-amgen-com/corlanor/corlanor\\_pi.pdf](https://www.pi.amgen.com/~media/amgen/repositorysites/pi-amgen-com/corlanor/corlanor_pi.pdf). Accessed November 19, 2019.
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**ii Egrifta SV References:**

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2. Clinical Pharmacology. <http://www.clinicalpharmacology-ip.com/Default.aspx>. Accessed September 6, 2019.
3. Treatment of HIV-associated lipodystrophy. (2020). UpToDate. [https://www.uptodate.com/contents/treatment-of-hiv-associated-lipodystrophy?search=egrifta&source=search\\_result&selectedTitle=2~3&usage\\_type=default&display\\_rank=1](https://www.uptodate.com/contents/treatment-of-hiv-associated-lipodystrophy?search=egrifta&source=search_result&selectedTitle=2~3&usage_type=default&display_rank=1). Accessed March 21, 2022.
4. Stanley T, Falutz J, Marsolais C, et al. Reduction in visceral adiposity is associated with an improved metabolic profile in HIV-infected patients receiving tesamorelin. Clin Infect Dis. 2012 Jun;54(11):1642-51. Accessed September 12, 2019
5. Clinical Review Report: Tesamorelin (Egrifta) [Internet]. Ottawa (ON): Canadian Agency for Drugs and Technologies in Health; 2016 Aug. <https://www.ncbi.nlm.nih.gov/books/NBK539131/> Accessed September 6, 2019

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